



DEPARTMENT OF JUSTICE  
DRUG ENFORCEMENT ADMINISTRATION  
MANUFACTURER OF CONTROLLED SUBSTANCES  
NOTICE OF APPLICATION  
PHARMAGRA LABS, INC.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 3, 2012, Pharmagra Labs, Inc., 158 McLean Road, Brevard, North Carolina 28712, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed substance for analytical research and clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement

Administration, Office of Diversion Control, Federal  
Register Representative (ODL), 8701 Morrisette Drive,  
Springfield, Virginia 22152; and must be filed no later  
than [INSERT DATE 60 DAYS FROM DATE OF PUBLICATION].

Joseph T. Rannazzisi  
Deputy Assistant Administrator  
Office of Diversion Control  
Drug Enforcement Administration

DATED: February 8, 2013

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